UNIVERSITÉ LAVAL

Faculté de médecine
Département de pédiatrie

Publications – Département de pédiatrie – 2021

1. ABHD16A deficiency causes a complicated form of hereditary spastic paraplegia associated with intellectual disability and cerebral anomalies. Lemire G, Ito YA, Marshall AE, **Chrestian N**, Stanley V, Brady L, Tarnopolsky M, Curry CJ, Hartley T, Mears W, Derksen A, Rioux N, Laflamme N, Hutchison HT, Pais LS, Zaki MS, Sultan T, Dane AD; Care4Rare Canada Consortium, Gleeson JG, Vaz FM, Kernohan KD, Bernard G, Boycott KM. Am J Hum Genet. 2021 Oct 7;108(10):2017-2023. doi: 10.1016/j.ajhg.2021.09.005. Epub 2021 Sep 28.PMID: 34587489

2. A Collaborative Model to Implement Flexible, Accessible and Efficient Oncogenetic Services for Hereditary Breast and Ovarian Cancer: The C-MOnGene Study Julie Lapointe, Michel Dorval, Jocelyne Chiquette, Yann Joly, Jason Robert Guertin , Maude Laberge, **Jean Gekas** , Johanne Hébert , Marie-Pascale Pomey, Tania Cruz-Marino, Omar Touhami, Arnaud Blanchet Saint-Pierre, Sylvain Gagnon, Karine Bouchard, Josée Rhéaume, Karine Boisvert, Claire Brousseau, Lysanne Castonguay, Sylvain Fortier, Isabelle Gosselin, Philippe Lachapelle, Sabrina Lavoie, Brigitte Poirier, Marie-Claude Renaud, Maria-Gabriela Ruizmangas, Alexandra Sebastianelli, Stéphane Roy, Madeleine Côté, Marie-Michelle Racine, Marie-Claude Roy, Nathalie Côté, Carmen Brisson, Nelson Charette, Valérie Faucher, Josianne Leblanc, Marie-Ève Dubeau, Marie Plante, Christine Desbiens, Martin Beaumont, Jacques Simard, Hermann Nabi Cancers (Basel) 2021 May 31;13(11):2729 doi: 10.3390/cancers13112729.

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